one of claims 1-13, the conjugate according to claim 14, or the pharmaceutical composition of claim 15 in an effective amount to said cell.

- 17. A method for treating or preventing a disease comprising administering a therapeutically or prophylactically effective amount of an oligonucleotide of any one of claims 1-13 or the conjugate according to claim 14 or the pharmaceutical composition of claim 15 to a subject suffering from or susceptible to the disease.
- 18. The method of claim 17, wherein the disease is a neurological disorder, such as a neurological disorder selected from the group consisting of Amyotrophic Lateral Sclerosis (ALS), Frontotemporal Dementia (FTD), tauopathy (such as primary tauopathy), frontotemporal dementia with parkinsonism (FTDP-17), frontotemporal lobar dementia (FTLD-TDP), Huntington's disease, Creutzfeld-Jacob disease, and spinomuscular atrophy, motor neuron disease, Tauopathy, Alzheimer's disease, and Welander distal myopathy.
- 19. The oligonucleotide of any one of claims 1-13 or the conjugate according to claim 14 or the pharmaceutical composition of claim 15 for use in medicine.
- 20. The oligonucleotide of any one of claims 1-13 or the conjugate according to claim 14 or the pharmaceutical composition of claim 15 for use in the treatment or prevention of a neurological disorder, such as a neurological disorder selected from the group consisting of Amyotrophic Lateral Sclerosis (ALS), Frontotemporal Dementia (FTD),

- tauopathy (such as primary tauopathy), frontotemporal dementia with parkinsonism (FTDP-17), frontotemporal lobar dementia (FTLD-TDP), Huntington's disease, Creutzfeld-Jacob disease, and spinomuscular atrophy, motor neuron disease, Tauopathy, Alzheimer's disease, and Welander distal myopathy.
- 21. Use of the oligonucleotide of claims 1-13 or the conjugate according to claim 14 or the pharmaceutical composition of claim 15, for the preparation of a medicament for treatment or prevention of a neurological disorder, such as a neurological disorder selected from the group consisting of Amyotrophic Lateral Sclerosis (ALS), Frontotemporal Dementia (FTD), frontotemporal dementia with parkinsonism (FTDP-17), frontotemporal lobar dementia (FTLD-TDP), tauopathy (such as primary tauopathy), Huntington's disease, Creutzfeld-Jacob disease, and spinomuscular atrophy, motor neuron disease, Tauopathy, Alzheimer's disease, and Welander distal myopathy.
- **22**. The Use of method according to any one of claims **17-21**, wherein the neurological disorder is Amyotrophic Lateral Sclerosis (ALS).
- 23. The Use of method according to any one of claims 17-21, wherein the neurological disorder is a tauopathy, such as a primary tauopathy.
- **24**. The Use of method according to any one of claims **17-21**, wherein the neurological disorder is frontotemporal lobar dementia (FTLD-TDP).

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